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(21) International Application Number: PCT/US94/14017 (22) International Filing Date: 5 December 1994 (05.12.94) (30) Priority Data: 08/170,460 20 December 1993 (20.12.93) US (71) Applicant: FERRO CORPORATION [US/US]; 1000 Lakeside Avenue, Cleveland, OH 44114-1183 (US). (72) Inventors: WIMOLKIATISAK, Surachai; 10811 Gate Post Road, Strongsville, OH 44136 (US). HAMMOND, Dennis, L.; 4278 Hart Road, Richfield, OH 44286 (US). SCHEIBELHOFFER, Anthony, S.; 2986 Wilbanks Drive, Norton, OH 44203 (US). CARLSON, Allen, W.; 160 North Maple Avenue, Ridgewood, NJ 07450 (US). ALI, Mir, L.; 524 Wesley Avenue, Bryan, OH 43506 (US). (74) Agent: CLARK, Kenneth, A.; Rankin, Hill, Lewis & Clark, Suite 600, 925 Euclid Avenue, Cleveland, OH 44115 (US).	(81) Designated States: AM, AT, AU, BB, BG, BR, BY, CA, CH, CN, CZ, DE, DK, ES, FI, GB, GE, HU, JP, KE, KG, KP, KR, KZ, LK, LT, LU, LV, MD, MG, MN, MW, NL, NO, NZ, PL, PT, RO, RU, SD, SE, SI, SK, TJ, TT, UA, UZ, VN, European patent (AT, BE, CH, DE, DK, ES, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE), OAPI patent (BF, BJ, CF, CG, CI, CM, GA, GN, ML, MR, NE, SN, TD, TG), ARIPO patent (KE, MW, SD, SZ). Published <i>With international search report.</i>	
(54) Title: VINYL HALIDE POLYMER COLOR CONCENTRATE (57) Abstract <p>The present invention provides a new and improved method of producing a dustless vinyl halide polymer color concentrate that does not require the formation of a hot melt or the use of wax. The color concentrate comprises a multitude of fine particles which each comprise a vinyl halide polymer resin core and a colorant material shell. The method includes the steps of providing a vinyl halide polymer, providing a colorant material, providing a compatibilizer having a solubility parameter of from about 7 to about 12 (cal/ml)^{1/2}, and mixing the vinyl halide polymer, compatibilizer and colorant material together to yield the dustless concentrate in the form of a vinyl halide resin core surrounded by a shell of the compatibilizer and the colorant material.</p>		

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Title: VINYL HALIDE POLYMER COLOR CONCENTRATE

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Field Of Invention

The present invention concerns a color concentrate for use in coloring plastics. More particularly, the present invention concerns a dustless solid color concentrate comprising a vinyl halide polymer resin core and a colorant material additive shell.

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Background

The traditional method for incorporating dry colors and additives into a thermoplastic polymer to produce a color concentrate involves blending the polymer with the color and additives in a melt, the melt being formed in an extruder or a batch type mixer such as a Banbury mixer. The disadvantages associated with these techniques include appreciable downtime for cleaning the equipment, high-energy costs for running the equipment, and long production cycles. Additionally, with respect to some polymer resins, the traditional method of

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5 processing can lead to the degradation of the polymer and/or the creation of an undesirable heat history.

Various attempts have been made to avoid using traditional methods in the production of color concentrates. One of these methods is disclosed in Aubrey et al., U.S. Patent No. 3,591,409. Aubrey et al. discloses a method for preparing coated resin granules wherein a mixture of resin granules, a wax and a particulate material is subjected to high-intensity blending at elevated temperatures. The Aubrey et al. process is applicable to a variety of thermoplastic resins including polyolefins, vinylidene polymers, polyamides and other polymers. One disadvantage associated with the Aubrey et al. process is that the wax employed has to be melted for the process to work. This makes it difficult to determine how much wax is required in order to coat the resin. Additionally, the wax may be considered in some applications to be a costly additive and/or an undesirable additive. For example, waxes can melt in the screw of an extruder or injection molding machine and this can cause lubrication of the screw and loss of production efficiency or rate.

25 Another method for incorporating color and additives into a polymer resin is described in Chatterjee et al., European Patent Publication No. 0,471,463A2. Chatterjee et al. discloses the use of high-speed impingement to incorporate color and additives into a polymer resin. This process, like

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5 the process disclosed in Aubrey et al., may be used to process
a variety of polymers. A disadvantage associated with this
process is that it requires the melting of the additive and/or
the surface of the polymer resin to promote the incorporation
of the additive into the polymer resin. If the processing
10 temperature is higher than the optimum temperature, the entire
polymer resin particle can melt, resulting in a large molten
mass or a large proportion of agglomerates instead of a
uniform powder size.

Another method for incorporating color and additives into
15 a polymer resin is described in Pennie et al., U.S. Patent No.
4,375,520. The method comprises the mixing in a Papenmeir
mixer of a liquid polymeric substance and a low molecular
weight polymer having a melting point in the range of about
95°C to 105°C. Pennie et al. is specific to the use of a
20 polymer having a melting point of about 95°C to 105°C and the
process is conducted at or near the melting point of the
polymer. Unfortunately, Pennie et al. does not provide any
insight into how to process a primarily amorphous polymer
resin that does not display a melting point, such as
25 poly(vinyl chloride) (PVC).

Summary Of Invention

The present invention provides a new and improved method
of producing a dustless vinyl halide polymer color concentrate
that does not require the formation of a hot melt or the use

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5 of wax. The color concentrate comprises a multitude of fine particles which each comprise a vinyl halide polymer resin core and a colorant material shell. The method includes the steps of providing a vinyl halide polymer, providing a colorant material, providing a compatibilizer having a solubility parameter of from about 7 to about 12 (cal/ml)^{1/2}, and mixing the vinyl halide polymer, compatibilizer and colorant material together to yield the dustless concentrate in the form of a vinyl halide resin core surrounded by a shell of the compatibilizer and the colorant material.

10 In a preferred embodiment the vinyl halide polymer comprises poly(vinyl chloride) (PVC) and the compatibilizer comprises epoxidized soybean oil.

15 These and other aspects of the present invention will become clear to those skilled in the art upon the reading and understanding of the specification and the claims below.

Detailed Description

20 Color concentrates made in accordance with the principles of the present invention yield various advantages. These advantages include for example: (i) the production of a non-dusting, free flowing, uniform particle size concentrate; (ii) the production of a concentrate that provides good dispersion when added to the thermoplastic being colored; (iii) the absence of melt extrusion or melt formation during production of the concentrate which can create an undesirable heat

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5 history; (iv) the absence of costly waxes that may cause
processing problems or detrimental properties; (v) batch size
flexibility; (vi) no screw slippage when the concentrate is
added to an extruder; (vii) faster color changes; (viii) the
absence of additives or other materials that can be
10 detrimental to the properties of the vinyl halide polymer
resin; and (ix) good yields during the production of the
concentrate.

Generally, the method comprises the steps of providing a
vinyl halide polymer resin, providing a compatibilizer having
15 a solubility parameter of from about 7 to about $12(\text{cal/ml})^{1/2}$,
providing one or more colorant materials, and mixing the
polymer resin, colorant and compatibilizer in a mixing device
such as a Henschel mixer.

The vinyl halide polymer resin may comprise one or more
20 of the commercially available polymer resins. Such polymers
include, for example, poly(vinyl chloride) including
homopolymers and copolymers thereof, alloys and blends
comprising homopolymers and copolymers of poly(vinyl
chloride), poly(vinyl fluoride) including homopolymers and
25 copolymers thereof, poly(vinylidene dichloride) including
homopolymers and copolymers thereof and chlorinated poly(vinyl
chloride) including homopolymers and copolymers thereof. The
vinyl halide polymer resin preferably has an average particle
size in the range of about 70 microns to about 500 microns.

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5 More preferably, the vinyl halide polymer resin has a particle size in the range of about 150 microns to about 250 microns.

The compatibilizer is a material that is liquid below 150°F and capable of being absorbed by the pigments and vinyl halide polymer resin being utilized. The compatibilizer has a solubility parameter of from about 7 to about 12 (cal/ml)^{1/2}.
10 More preferably, the compatibilizer has a solubility parameter of from about 8.5 to about 10.5 (cal/ml)^{1/2}. Examples of preferred compatibilizers include epoxidized soybean oil, epoxidized tall oil, epoxidized linseed oil, ethoxylated sorbitan trioleate, sorbitan trioleate, tri-iso-octyl phosphate, octyl diphenyl phosphate, tolyl diphenyl phosphate, tritolyl phosphate, trixylyl phosphate, triphenyl phosphate, di-n-butyl phthalate, di-2-ethylhexyl phthalate, di-n-hexyl phthalate, di-iso-octyl phthalate, di-n-octyl phthalate, di-
15 isodecyl phthalate, di-tridecyl phthalate, n-Butyl benzyl phthalate, di-2-methoxyethyl phthalate, butyl butoxycarbonylmethyl phthalate, di-2-ethylhexyl adipate, iso-octyl isodecyl adipate, di-isodecyl adipate, di-2-ethylhexyl azelate, di-iso-octyl azelate, di-2-ethylhexyl sebacate, and
20 chlorinated paraffin extender. Preferably, when using poly(vinyl chloride) as the polymer resin, the compatibilizer comprises epoxidized soybean oil.

The colorant material may comprise any one or more of a number of commercially available colors or pigments. As used

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5 in this specification and claims below "colorant material"
means any conventional inorganic or organic pigment or organic
dyestuff. Such materials are described, for example, in Kirk-
Othmer Encyclopedia of Chemical Technology, Third Edition,
Vol. 6, pp. 597-617, which is incorporated herein by
10 reference. Examples of inorganic pigments include, for
example, titanium dioxide, iron oxide, zinc chromate, cadmium
sulfides, chromium oxides and sodium aluminum silicate
complexes. Examples of organic type pigments include azo and
diaz pigments, phthalocyanines, quinacridone pigments,
15 perylene pigments, isoindolinone, anthraquinones, thioindigo
and solvent dyes.

 Various types of additives may also be included in the
process. Such additives may include, for example, stabilizers
such as alkaline earth metal soaps and carboxylates such as
20 calcium benzoate, calcium octoate and calcium naphthenate,
Friedel-Crafts cation progenitor compounds such as zinc oxide,
zinc hydroxide, zinc carbonate, zinc acetate, zinc laurate,
zinc naphthenate, zinc stearate, zinc oleate, zinc 2-ethyl-
hexoate, cadmium oxide, cadmium hydroxide, cadmium carbonate,
25 cadmium acetate, cadmium laurate, cadmium naphtheneate,
cadmium stearate, cadmium oleate, cadmium 2-ethyl-hexoate,
calcium stearate, aluminum oxide, aluminum hydroxide, aluminum
carbonate, aluminum acetate, aluminum chlorolaurate, antimony
oxide, antimony hydroxide, antimony carbonate, antimony

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5 naphthenate, tin oxide, tin hydroxide, tin carbonate, tin
naphthenate, and tin 2-ethyl-hexanoate, aliphatic polyhydric
compounds such as trimethylolethane, trimethylolpropane,
tetramethylolcyclohexanol, pentaerythritol, dipentaerythritol,
and tripentaerythritol and tri-(2-hydroxyethyl) iso-cyanurate
10 (THEIC).

Various types of fillers and/or reinforcers may also be
included in the process. Such fillers and/or reinforcers
include, for example, CaCO_3 , talc, glass, clay and mica.

15 The portions of vinyl halide polymer resin, color and
compatibilizer can vary quite a bit. Generally, when
producing the concentrate, the mixture utilized comprises from
about 20 to about 60 weight percent vinyl halide polymer, from
about 5 to about 20 weight percent compatibilizer and from
about 25 to about 75 weight percent colorant material. The
20 mixture may comprise up to 15 weight percent additive.
Preferably, the mixture comprises from about 25 to about 55
weight percent vinyl halide polymer, from about 10 to about 15
weight percent compatibilizer, and from about 30 to about 70
weight percent colorant material.

25 Various high-intensity mixing devices may be utilized in
accordance with the principles of the present invention. Such
devices include, for example, Papenmeier mixers, Waring
blenders and Henschel mixers. Preferably, a Henschel mixer is
employed. Generally, the mixing device is operated at a tip

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5 speed of from about 10 to about 60 meters/second. Preferably,
the mixing device is operated at a tip speed of from about 20
to about 40 meters/second.

Generally, if an additive is employed, it is best when
producing the concentrate to first mix the additive with the
10 vinyl halide polymer resin for a brief period such as 10 to 20
seconds. The colorant material then can be added and mixed
for 10 to 20 seconds. The compatibilizer is then added and
mixed for 3 to 10 minutes. Of course, it will be appreciated
that these are merely general mixing times and the actual
15 mixing times will depend on such variables as, for example,
the size of the mixing container, the amount and type of
materials being mixed and the speed and configuration of the
mixing blade.

Care must be taken not to allow the mixture during the
20 mixing operation to exceed the degradation temperature of the
polymer resin (215°F for PVC). Thus, when working with PVC a
maximum of about 210°F should be attained. To obtain a free-
flowing concentrate, the mixing should be conducted until a
temperature of from about 140°F to about 180°F is reached, and
25 preferably from about 150°F to about 180°F is reached.
Excessively high temperature should be avoided because it can
cause the vinyl halide polymer resin to degrade. If during
mixing smearing should result, metal stearates, such as
calcium stearate, can be added to the mixture to prevent

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5 smearing. Generally, such additions are made in the range of .5% to about 1%. Addition of metal stearates to the mixer may result in the formation of a fragile ball. However, such ball will readily break up into free-flowing particles with a gentle tap.

10 After mixing the mixture is generally passed through a sieve in order to remove oversize aggregates. Generally, a yield of over 90% by weight is attained. After sieving the mixture comprises a non-dusting, free-flowing, uniform particle size concentrate. The concentrate generally
15 comprises a multitude of particles having an average particle size of from about 150 to about 850 microns. Each particle comprises a vinyl halide polymer resin core and an outer coating of compatibilizer, colorant materials and additives. The resin cores have a diameter of from about 70 microns to
20 about 500 microns.

 Any number of let-down resins can be used in connection with the color concentrates of the present invention. For example, the let-down resin can be either flexible or rigid PVC compounds or any other polymers compatible with the vinyl
25 halide resin used to produce the color concentrate. The color concentrates of the present invention are used in the same manner as conventional concentrates (i.e., dry blending with the let-down resin followed by extrusion). Let-down ratios

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5 vary depending on the desired colors and applications. In general, the ratio ranges from about 10:1 to about 100:1.

During manufacture the ultimate color of the color concentrate can be easily controlled or corrected. More particularly, before adding the compatibilizer one can conduct a color check on the colorant material and vinyl halide polymer resin using any one of a number of conventional techniques, such as, for example, melt mixing with the let-down resin. If the color of the melt is found to be unacceptable, additional colorant materials can be added during manufacture prior to the addition of the compatibilizer so as to adjust the color as required.

The following examples will serve to illustrate the novel features and advantages of the present invention. While these examples will show one skilled in the art how to operate within the scope of this invention, they are not to serve as a limitation on the scope of the invention for such scope is only defined in the claims below.

Example I

The following components were charged into a FM150 Henschel mixer having a 150 litre mixing chamber. The mixer was operated at a tip speed of 20 meters/second using a variant-type mixing blade.

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	<u>Component</u>	<u>Weight %</u>	<u>Weight</u>
5	GEON 110 x 334 PVC	37	35.15 lbs.
	Ferro Corporation V6797 mixed metal oxide pigment	50	47.5 lbs.
	Calcium stearate	3	2.85 lbs.
10	Ferro Corporation THERM-CHEK 840 stabilizer	.5	.475 lbs.
	Epoxidized soybean oil	9.5	9.025 lbs.

15 The PVC and the THERM-CHEK 840 stabilizer are mixed separately for 15 seconds. The calcium stearate and mixed metal pigment are then added and mixed for 15 seconds. The epoxidized soybean oil is then gradually added and mixed for five minutes yielding a mixture of free-flowing fine particles. The fine particles are then passed through a 20 mesh screen yielding about 2% oversized particles that do not pass through the screen. The sieved particles are ready for use as a color concentrate. The oversize particles can be ground and used in subsequent batches without causing any detrimental effects.

Example II

25 The following components were charged into a FM150 Henschel mixer having a 150 litre mixing chamber. The mixer was operated at a tip speed of 20 meters/second using a variant-type mixing blade.

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5	<u>Component</u>	<u>Weight%</u>	<u>Weight</u>
	Carbon black	1.6	1.52 lbs.
	Ethylene bis-stearamide	2	1.9 lbs.
	Calcium carbonate	30	28.5 lbs.
	Calcium stearate	2.5	2.38 lbs.
10	Vygen 310 PVC	55.4	52.6 lbs.
	Epoxidized soybean oil	8	7.6 lbs.
	Ferro Corporation THERM-CHEK 840 stabilizer	.5	.48 lbs.

15 The PVC and the THERM-CHEK 840 stabilizer are mixed separately for 15 seconds. 1.9 pounds of the calcium stearate, the carbon black, the ethylene bis-stearamide and the calcium carbonate are then gradually added and mixed for 15 seconds. The epoxidized soybean oil is then added and mixed for about 5 minutes. Finally, .48 pounds of the calcium

20 stearate is added and mixed for 30 seconds to eliminate smearing. The calcium stearate added in the final step coats each of the fine particles that are produced. The fine particles are then passed through a 20 mesh screen yielding about 2% oversize particles that do not pass through the

25 screen. The sieved particles are ready for use as a color concentrate. The oversized particles can be ground and used in subsequent batches without causing any detrimental effects.

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Example III

The color concentrate made as described in Example I was used at a 33:1 let-down ratio with a white rigid PVC compound. The two materials were dry blended and melt extruded to make a 1" diameter gray PVC pipe. The materials extruded well and the pipe has good appearance with good pigment dispersion.

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Example IV

The color concentrate made as described in Example II was used at a 20:1 let-down ratio with a white rigid PVC compound. The two materials were dry blended and melt mixed on a two-roll mill to make a gray PVC sheet. The materials processed well and the sheet had good appearance with good pigment dispersion.

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Although the invention has been shown and described above with respect to specific embodiments, it is obvious that equivalent alterations and modifications will occur to others skilled in the art upon reading and understanding the specification. The present invention includes all such equivalent alterations and modifications, and is limited only by the scope of the following claims.

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5 **Claims:**

1. A method of producing a dustless, free-flowing vinyl halide polymer color concentrate comprising a multitude of fine particles each comprising a vinyl halide polymer resin core and a colorant material shell, said method comprising the steps of:

10

A. providing a vinyl halide polymer;

B. providing a compatibilizer having a solubility parameter of from about 7 to about 12 (cal/ml)^{1/2};

C. providing a colorant material;

15

D. mixing such vinyl halide polymer, such compatibilizer and such colorant in a container to provide a mix and form such color concentrate comprising a multitude of fine particles.

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2. A method as set forth in claim 1 wherein such vinyl halide polymer comprises poly(vinyl chloride).

25

3. A method as set forth in claim 1 wherein such compatibilizer is selected from the group consisting essentially of epoxidized soybean oil, epoxidized tall oil, epoxidized linseed oil, ethoxylated sorbitan trioleate, or sorbitan trioleate, and said mixing step D is conducted using a high intensity mixer with a tip speed of from about 20 to about 40 meters/second.

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5 4. A method as set forth in claim 1 wherein such container is maintained at a temperature below the degradation and melting temperatures of such vinyl halide polymer.

5. A method as set forth in claim 1 wherein said mixing step D is conducted using a Henschel mixer.

10 6. A method as set forth in claim 1 including the step of providing an additive material and mixing such additive material with such vinyl halide polymer, such compatibilizer and such colorant during said step D.

15 7. A method as set forth in claim 1 wherein during said step D such vinyl halide polymer, such compatibilizer and such colorant material are mixed at a temperature of from about 140°F to about 180°F.

20 8. A method as set forth in claim 1 wherein during said step D such vinyl halide polymer, such compatibilizer and such colorant material are mixed at a maximum of about 210°F.

25 9. A method as set forth in claim 1 wherein the mix of said step D comprises from about 20 to about 60 weight percent vinyl halide polymer, from about 5 to about 20 weight percent compatibilizer, and from about 25 to about 75 weight percent colorant material.

10. A method as set forth in claim 1 wherein the mix of said step D comprises from about 25 to about 55 weight percent vinyl halide polymer, from about 10 to about 15 weight percent

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5 compatibilizer, and from about 30 to about 70 weight percent
colorant material.

11. A method as set forth in claim 1 wherein such vinyl
halide polymer of said step A has an average particle size of
from about 70 microns to about 500 microns.

10 12. A method as set forth in claim 1 wherein such vinyl
halide polymer of said step A has an average particle size of
from about 250 microns to about 500 microns.

13. A method as set forth in claim 1 wherein such
compatibilizer of said step B has a solubility parameter of
15 from about 8.5 to about 10.5 (cal/ml)^{1/2}.

14. A method as set forth in claim 1 including the step
of providing an additive and mixing such additive with such
mix during said step D.

15. A method as set forth in claim 14 wherein such
20 additive comprises calcium stearate.

16. A method as set forth in claim 1 wherein said
colorant material comprises an inorganic pigment.

17. A method as set forth in claim 1 wherein said
colorant material comprises an organic pigment.

25 18. A method as set forth in claim 1 wherein said color
concentrate comprises a multitude of free-flowing particles
which display an average particle size of from about 150
microns to about 850 microns, said particles comprising a

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5 vinyl halide polymer core and an outer coating of
compatibilizer and colorant material.

19. A method as set forth in claim 1 wherein during said
step D a color check is conducted prior to the addition of
such compatibilizer.

10 20. A dustless free-flowing poly(vinyl chloride) based
color concentrate comprising a multitude of particles which
display an average particle size of from about 150 microns to
about 850 microns, said particles comprising a poly(vinyl
chloride) core and an outer coating of compatibilizer and
15 colorant material.

21. A color concentrate as set forth in claim 20 wherein
said cores of said particles have a diameter of from about 70
microns to about 500 microns and said color concentrate is
substantially free of wax.

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INTERNATIONAL SEARCH REPORT

International application No.
PCT/US94/14017

A. CLASSIFICATION OF SUBJECT MATTER

IPC(6) :B05D 1/00, 3/12
US CL :427/212,218,220,222; 428/403
According to International Patent Classification (IPC) or to both national classification and IPC

B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)

U.S. : 427/212,218,220,222; 428/403

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched
NONE

Electronic data base consulted during the international search (name of data base and, where practicable, search terms used)
NONE

C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category*	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
X --- Y	US, A, 3,819,560 (KEHR) 25 JUNE 1974, COL. 2, LINES 23-62 AND COL. 3, LINES 5-8 AND COL. 3, LINES 55-65 AND COL. 8, LINES 13-46.	1, 2, 4, 5, 11-14, 16-18 ----- 1-19
Y	US, A, 3,591,409 (AUBREY ET AL.) 06 JULY 1971, ABSTRACT AND COL. 2, LINES 16-54 AND COL. 3, LINES 33-74 AND COL. 4, LINES 47-62 AND COL. 5, LINES 10-30 AND COL. 6, LINES 27-31.	1-19
Y	EP, A, 0 471 463 (CHATTERJEE ET AL.) 19 FEBRUARY 1992, ABSTRACT AND COL. 3, LINES 9-49.	1-19
A	US, A, 5,139,817 (ABE ET AL.) 18 AUGUST 1992, ENTIRE REFERENCE	1-19

☒ Further documents are listed in the continuation of Box C. ☐ See patent family annex.

* Special categories of cited documents:	*T	later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention
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P document published prior to the international filing date but later than the priority date claimed		

Date of the actual completion of the international search
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09 MAR 1995

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C (Continuation). DOCUMENTS CONSIDERED TO BE RELEVANT		
Category*	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
A	US, A, 5,264,280 (CHUNDURY ET AL.) 23 NOVEMBER 1993, ENTIRE REFERENCE	1-19
A	US, A, 5,066,422 (FELTER ET AL.) 19 NOVEMBER 1991, ENTIRE REFERENCE	1-19
A	US, A, 4,375,520 (PENNIE ET AL.) 01 MARCH 1983, ENTIRE REFERENCE	1-19
A	US, A, 4,495,128 (STOFFELSMA) 22 JANUARY 1985, ENTIRE REFERENCE	1-19
A	CA, A, 2,006,350 (BES ET AL.) 27 JUNE 1990, ENTIRE REFERENCE	1-19